

FILE 'REGISTRY' ENTERED AT 15:30:18 ON 02 MAR 2009  
EXP LINOLEIC/CN  
L1 1 S E4

FILE 'HCAPLUS' ENTERED AT 15:30:46 ON 02 MAR 2009  
L2 64068 S L1 OR LINOLEIC OR (VITAMIN F) OR (OMEGA-6)  
L3 39402 S CYCLODEXTRIN  
L4 189 S L2 AND L3  
L5 106 S L4 AND (PY<2003 OR AY<2003 OR PRY<2003)

FILE 'HCAPLUS' ENTERED AT 15:35:28 ON 02 MAR 2009

FILE 'REGISTRY' ENTERED AT 15:35:37 ON 02 MAR 2009  
E A CYCLODEXTRIN/CN

FILE 'HCAPLUS' ENTERED AT 15:35:37 ON 02 MAR 2009

FILE 'REGISTRY' ENTERED AT 15:35:47 ON 02 MAR 2009  
E ACYCLODEXTRIN/CN

FILE 'HCAPLUS' ENTERED AT 15:35:48 ON 02 MAR 2009

FILE 'REGISTRY' ENTERED AT 15:36:04 ON 02 MAR 2009  
E ALPHA CYCLODEXTRIN/CN

FILE 'HCAPLUS' ENTERED AT 15:36:05 ON 02 MAR 2009  
L6 7087 S ALPHA CYCLODEXTRIN  
L7 38 S L2 AND L6  
L8 23 S L7 AND (PY<2003 OR AY<2003 OR PRY<2003)

=> file registry  
COST IN U.S. DOLLARS  
FULL ESTIMATED COST

SINCE FILE	TOTAL
ENTRY	SESSION
0.22	0.22

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Property values tagged with IC are from the ZIC/VINITI data file  
provided by InfoChem.

STRUCTURE FILE UPDATES: 27 FEB 2009 HIGHEST RN 1113101-98-6  
DICTIONARY FILE UPDATES: 27 FEB 2009 HIGHEST RN 1113101-98-6

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH January 9, 2009.

Please note that search-term pricing does apply when  
conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and  
predicted properties as well as tags indicating availability of  
experimental property data in the original document. For information  
on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

=> exp linoleic/cn

E1	1	LINOLEATE ISOMERASE (LACTOBACILLUS PLANTARUM STRAIN AS1.555) /CN
E2	1	LINOLEATE PEROXYL RADICAL/CN
E3	0 -->	LINOLEIC/CN
E4	1	LINOLEIC ACID/CN
E5	1	LINOLEIC ACID (D(-)-), (2,2-DIMETHYL-1,3-DIOXOLAN-4-YL)METHY L ESTER/CN
E6	1	LINOLEIC ACID (L(-)-), 2-HYDROXY-3-(TRILYLOXY)PROPYL ESTER/C N
E7	1	LINOLEIC ACID $\Omega$ -6 LIPOXYGENASE/CN
E8	1	LINOLEIC ACID 1-(2-NAPHTHYL)ETHYL ESTER/CN
E9	1	LINOLEIC ACID 1-NAPHTHYLMETHYL ESTER/CN
E10	1	LINOLEIC ACID 10-HYDROPEROXIDE/CN
E11	1	LINOLEIC ACID 12-HYDROPEROXIDE/CN
E12	1	LINOLEIC ACID 13(S)-HYDROPEROXIDE/CN

=> s e4

L1 1 "LINOLEIC ACID"/CN

=> file hcaplus  
COST IN U.S. DOLLARS  
FULL ESTIMATED COST

SINCE FILE	TOTAL
ENTRY	SESSION
5.83	6.05

FILE 'HCAPLUS' ENTERED AT 15:30:46 ON 02 MAR 2009  
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FILE COVERS 1907 - 2 Mar 2009 VOL 150 ISS 10  
FILE LAST UPDATED: 1 Mar 2009 (20090301/ED)

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

CAS Information Use Policies apply and are available at:

<http://www.cas.org/legal/infopolicy.html>

This file contains CAS Registry Numbers for easy and accurate substance identification.

```
=> s l1 or linoleic or (vitamin F) or (omega-6)
      43084 L1
      48263 LINOLEIC
      217991 VITAMIN
      671725 F
      347 VITAMIN F
          (VITAMIN(W)F)
      201652 OMEGA
      4246472 6
      6147 OMEGA-6
          (OMEGA(W)6)
L2      64068 L1 OR LINOLEIC OR (VITAMIN F) OR (OMEGA-6)
```

```
=> s cyclodextrin
L3      39402 CYCLODEXTRIN
```

```
=> log hold
COST IN U.S. DOLLARS                SINCE FILE      TOTAL
                                     ENTRY      SESSION
FULL ESTIMATED COST                2.85      8.90
```

SESSION WILL BE HELD FOR 120 MINUTES  
STN INTERNATIONAL SESSION SUSPENDED AT 15:31:19 ON 02 MAR 2009

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID:SSPTAEXO1623

PASSWORD:

```
* * * * * RECONNECTED TO STN INTERNATIONAL * * * * *
SESSION RESUMED IN FILE 'HCAPLUS' AT 15:34:46 ON 02 MAR 2009
FILE 'HCAPLUS' ENTERED AT 15:34:46 ON 02 MAR 2009
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COST IN U.S. DOLLARS	SINCE FILE	TOTAL
FULL ESTIMATED COST	ENTRY	SESSION
	2.85	8.90

=> d his

(FILE 'HOME' ENTERED AT 15:30:09 ON 02 MAR 2009)

FILE 'REGISTRY' ENTERED AT 15:30:18 ON 02 MAR 2009  
EXP LINOLEIC/CN

L1 1 S E4

FILE 'HCAPLUS' ENTERED AT 15:30:46 ON 02 MAR 2009

L2 64068 S L1 OR LINOLEIC OR (VITAMIN F) OR (OMEGA-6)

L3 39402 S CYCLODEXTRIN

=> s l2 and l3

L4 189 L2 AND L3

=> s l4 and (PY<2003 or AY<2003 or PRY<2003)

22983504 PY<2003

4504574 AY<2003

3973543 PRY<2003

L5 106 L4 AND (PY<2003 OR AY<2003 OR PRY<2003)

=> file hcaplus

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
FULL ESTIMATED COST	ENTRY	SESSION
	5.70	11.75

FILE 'HCAPLUS' ENTERED AT 15:35:28 ON 02 MAR 2009

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FILE COVERS 1907 - 2 Mar 2009 VOL 150 ISS 10

FILE LAST UPDATED: 1 Mar 2009 (20090301/ED)

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

CAS Information Use Policies apply and are available at:

<http://www.cas.org/legal/infopolicy.html>

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> exp  $\alpha$  cyclodextrin/cn

REGISTRY INITIATED  
Substance data EXPAND from CAS REGISTRY in progress...

E1	1	A CHAIN HEMOGLOBIN (BATHYRAJA EATONII)/CN
E2	1	A CRYSTALLIN (RATTUS RATTUS STRAIN SPRAGUE-DAWLEY CLON E PARL3 B-CHAIN)/CN
E3	0 -->	A CYCLODEXTRIN/CN
E4	1	A E-CATENIN (SUS SCROFA C-TERMINAL FRAGMENT)/CN
E5	1	A ENOLASE (ALLIGATOR MISSISSIPPIENSIS MUSCLE)/CN
E6	34	A ENOLASE (AMEIVA CHRYSOLAEMA FRAGMENT)/CN
E7	1	A ENOLASE (ASTERIAS RUBENS STRAIN ENO3 FRAGMENT)/CN
E8	1	A ENOLASE (CAIMAN CROCODYLUS FRAGMENT)/CN
E9	1	A ENOLASE (CEREBRATULUS STRAIN ENO6 FRAGMENT)/CN
E10	1	A ENOLASE (ECHIURUS ECHIURUS STRAIN ENO4 FRAGMENT)/CN
E11	2	A ENOLASE (EUMECES INEXPECTATUS FRAGMENT)/CN
E12	1	A ENOLASE (EUPREPIS PERROTETII STRAIN SPECIMEN-VOUCHER -FMNH-262227 FRAGMENT)/CN

=> exp  $\alpha$ cyclodextrin/cn  
REGISTRY INITIATED  
Substance data EXPAND from CAS REGISTRY in progress...

E1	1	AB-CRYSTALLIN (TRICHECHUS MANATUS GENE CRYAB N-TERMINA L FRAGMENT)/CN
E2	1	AC-CONOTOXIN PRXA/CN
E3	0 -->	ACYCLODEXTRIN/CN
E4	1	AD-GLOBIN (KALINOWASKI'S TINAMOU)/CN
E5	1	AD-GLOBIN (KOMODO DRAGON REDUCED)/CN
E6	1	AE-CATENIN (DANIO RERIO)/CN
E7	1	AE-CATENIN (HUMAN GENE CTNNA1)/CN
E8	1	AF-INTERFERON (HUMAN)/CN
E9	1	AG HESPERIDIN/CN
E10	1	AG HESPERIDIN H/CN
E11	1	AG HESPERIDIN PA/CN
E12	1	AG HESPERIDIN PS/CN

=> exp  $\alpha$ cyclodextrin/cn  
REGISTRY INITIATED  
Substance data EXPAND from CAS REGISTRY in progress...

E1	1	ALPHA CONOTOXIN QCAL-1 (CONUS QUERCINUS)/CN
E2	1	ALPHA CONOTOXIN QCAL-2 (CONUS QUERCINUS)/CN
E3	0 -->	ALPHA CYCLODEXTRIN/CN
E4	1	ALPHA D3/CN
E5	1	ALPHA DNA POLYMERASE (CRYPTOCOCCUS NEOFORMANS NEOFORMANS STR AIN JEC21)/CN
E6	1	ALPHA E1 ESTERASE (HAEMATOBIA IRRITANS STRAIN CAMP COOLEY-4/ 97 GENE AE1)/CN
E7	1	ALPHA E7 ESTERASE (HAEMATOBIA IRRITANS IRRITANS STRAIN CAMP COOLEY 4/97 CLONE HF41CC GENE AE7)/CN
E8	1	ALPHA E8 ESTERASE (HAEMATOBIA IRRITANS STRAIN CAMP COOLEY-4/ 97 GENE AE8)/CN
E9	3	ALPHA ENDOSULFINE (HUMAN)/CN

E10 1 ALPHA ENOLASE (HUMAN CLONE 23942)/CN  
 E11 1 ALPHA ENOLASE LIKE 1 (HUMAN GENE ENO1L1)/CN  
 E12 1 ALPHA ENOLASE/TAU-CRYSTALLIN (FICEDULA HYPOLEUCA ISOLATE OS3 )/CN

=> s alpha cyclodextrin  
 1829388 ALPHA  
 39402 CYCLODEXTRIN  
 L6 7087 ALPHA CYCLODEXTRIN  
 (ALPHA(W)CYCLODEXTRIN)

=> s 12 and 16  
 L7 38 L2 AND L6

=> s 17 and (PY<2003 or AY<2003 or PRY<2003)  
 22983504 PY<2003  
 4504574 AY<2003  
 3973543 PRY<2003  
 L8 23 L7 AND (PY<2003 OR AY<2003 OR PRY<2003)

=> d 18 1-23 ti abs bib

L8 ANSWER 1 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN  
 TI Cosmetic composition comprising a complex of cyclodextrin and vitamin F  
 AB The invention concerns cosmetic and dermatol. compns. that contain complexes of vitamin F with  $\alpha$ ,  $\beta$ , or  $\gamma$ -cyclodextrin. Addnl. substances in the formulations are: silicone oils, moisturizers, skin care substances, gelation agents, bactericides, antioxidants, sunscreens, emulsifiers, pigments, tanning agents, etc. Thus 0.1 mol  $\alpha$  -cyclodextrin was mixed with 100 g water; 0.1 mol linolic acid was added, homogenized and stirred for 30 h at RT and for 8 h at 70°C; the product was dispersed in water, filtered, washed and dried under vacuum. A composition contained

(weight/weight%):.  
 alpha.-cyclodextrin-linolic acid complex 4.0;  
 $\gamma$ -cyclodextrin- $\alpha$ -tocopherol complex 1.5; octyl palmitate 2.5;  
 octyl stearate 3.5; polyglycerol-2 sesquiosostearate 2.0; cyclomethicone, dimethiconol 3.0; lauryl dimethicone 2.0; octyl dimethicone ethoxy glycoside, cyclomethicone 12.0; titanium dioxide 5.0; polymethylsilsesquioxane 1.0; zinc oxide 2.0; glycerin 2.0; methylparaben 0.1; sodium chloride 0.4; water 59.0.

AN 2004:402912 HCAPLUS <<LOGINID::20090302>>

DN 140:412001

TI Cosmetic composition comprising a complex of cyclodextrin and vitamin F

IN Regiert, Marlies; Kupka, Michaela

PA Wacker-Chemie GmbH, Germany

SO Eur. Pat. Appl., 17 pp.

CODEN: EPXXDW

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1419761	A1	20040519	EP 2003-26137	20031113 <--
	EP 1419761	B1	20051019		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	DE 10253042	A1	20040603	DE 2002-10253042	20021114 <--

KR 2004042827	A	20040520	KR 2003-77579	20031104 <--
US 20040096413	A1	20040520	US 2003-712703	20031112 <--
JP 2004161775	A	20040610	JP 2003-385675	20031114 <--
PRAI DE 2002-10253042	A	20021114	<--	

RE.CNT 12      THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 2 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN  
 TI Health-care wine of conjugated linoleic acid or its derivative  
 AB The health-care wine is composed of wine 1,000, antioxidant 0.01-20, conjugated linoleic acid or its derivative 0.1-100, taste improver 0.001-100, surfactant 0.001-20, and adjuvant 0.001-10 part. The drinkable wine is the distilled wine, grape wine, yellow rice wine, etc. The conjugated linoleic acid is 9,11-conjugated linoleic acid, 8,10-conjugated linoleic acid, 8,10-conjugated linoleic acid, and 11,13-conjugated linoleic acid. The conjugated linoleic acid derivative is inclusion complex of cyclodextrin or its derivative, C1-8 alkyl conjugated linoleate, ethylene bis(conjugated linoleate), glycerol mono- conjugated linoleate, glycerol di(conjugated linoleate), glycerol tri(conjugated linoleate), vitamin E conjugated linoleate, etc. The taste improver is sucrose, glucose, fructose, maltose, etc. The antioxidant is vitamin C, vitamin E, isoascorbic acid, etc. The surfactant is Span series, Tween series, sucrose ester, etc. The adjuvant is sucrose octa(acetate), hydroxypropyl starch, Na alginate, etc.

AN 2004:167321 HCAPLUS <<LOGINID::20090302>>

DN 140:198487

TI Health-care wine of conjugated linoleic acid or its derivative

IN Wumanjiang, Aili; Zhang, Yagang; Wen, Bin; Fan, Li; Ma, Li; Nu'ermaimaiti

PA Xinjiang Institute of Chemistry, Chinese Academy of Sciences, Peop. Rep. China

SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 14 pp.

CODEN: CNXXEV

DT Patent

LA Chinese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	---	-----	-----	-----
PI	CN 1371985	A	20021002	CN 2002-102288	20020205 <--
	CN 1200089	C	20050504		
PRAI	CN 2002-102288		20020205	<--	

L8 ANSWER 3 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Production method of cyclodextrin inclusion materials using marine or animal products

AB Title method comprise treatment of mixts. comprising lipophilic component-containing marine or animal products, starch, and lipid soluble solvents by addition of cyclodextrin synthetase. Thus, 5 g rice starch, 10 g salmon caviar, and 1 THU (based on 1 g starch) cyclodextrin synthetase were reacted in ethanol to give a cyclodextrin inclusion material showing good antioxidant property.

AN 2004:139298 HCAPLUS <<LOGINID::20090302>>

DN 140:182653

TI Production method of cyclodextrin inclusion materials using marine or animal products

IN Miwa, Shoji

PA Ishikawa Prefecture, Japan

SO Jpn. Kokai Tokkyo Koho, 13 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2004051866	A	20040219	JP 2002-213621	20020723 <--
	JP 4203578	B2	20090107		
PRAI	JP 2002-213621		20020723	<--	

L8 ANSWER 4 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Skin sanitizing compositions

AB The present invention relates to compns. and methods of sanitizing and moisturizing skin surfaces. A sanitizing and moisturizing gel contained EtOH 55, isopropanol 3, Biowax-754 0.4, Carbopol Ultrez-10 0.3, Carbowax PEG-200 0.26, propylene glycol 0.02, aminomethylpropanol 0.15, and perfume 0.1%, and water qs.

AN 2002:551533 HCAPLUS <<LOGINID::20090302>>

DN 137:114518

TI Skin sanitizing compositions

IN Sine, Mark Richard; Wei, Karl Shiqing; Jakubovic, David Andrew; Thomas, Cheyne P.; Dodd, Michael Thomas; Putman, Christopher Dean

PA The Procter & Gamble Company, USA

SO U.S., 14 pp., Cont. of U.S. Ser. No. 321,291.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 6423329	B1	20020723	US 2000-504286	20000215 <--
PRAI	US 1999-249717	A2	19990212	<--	
	US 1999-120098P	P	19990216	<--	
	US 1999-321291	A2	19990527	<--	

RE.CNT 56 THERE ARE 56 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 5 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Inclusion complex of conjugated linoleic acid (CLA) with cyclodextrins

AB Conjugated linoleic acid (CLA) inclusion complexes with . alpha.-cyclodextrin ( $\alpha$ -CD),  $\beta$ -cyclodextrin ( $\beta$ -CD), and  $\gamma$ -cyclodextrin ( $\gamma$ -CD) (designated CLA/CDs inclusion complexes) were prepared to determine the mole ratio of CLA complexed with CDs and the oxidative stability of CLA in the CLA/CDs inclusion complexes. When measured by GC, <sup>1</sup>H NMR, and T1 value analyses, 1 mol of CLA was complexed with 5 mol of  $\alpha$ -CD, 4 mol of  $\beta$ -CD, and 2 mol of  $\gamma$ -CD. The oxidation of CLA induced at 35° for 80 h was completely prevented by the formation of CLA/CDs inclusion complexes.

AN 2002:259433 HCAPLUS <<LOGINID::20090302>>

DN 137:19586

TI Inclusion complex of conjugated linoleic acid (CLA) with cyclodextrins

AU Park, Cherl W.; Kim, Seck J.; Park, Sook J.; Kim, Jeong H.; Kim, Jung K.; Park, Gu B.; Kim, Jeong O.; Ha, Yeong L.

CS Division of Applied Life Sciences and Institute of Agriculture and Life Sciences Graduate School, Gyeongsang National University, Jinju, 660-701, S. Korea

SO Journal of Agricultural and Food Chemistry (2002), 50(10), 2977-2983

CODEN: JAFCAU; ISSN: 0021-8561

PB American Chemical Society

DT Journal

LA English



RE.CNT 31        THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8    ANSWER 6 OF 23    HCAPLUS    COPYRIGHT 2009 ACS on STN  
TI    Improvement of Oxidative Stability of Conjugated Linoleic Acid  
      (CLA) by Microencapsulation in Cyclodextrins  
AB    Oxidative stability of conjugated linoleic acid (CLA)  
      encapsulated in  $\alpha$ -,  $\beta$ -, and  $\gamma$ -cyclodextrins (designated  
      CLA/CDs microencapsules) was studied by measuring the headspace-oxygen  
      depletion in airtight serum bottles and by measuring the peroxide values  
      (POV). The rate of oxygen depletion was reduced from 41.0 (control) to  
      21.5, 2.1, 1.2, and 1.1  $\mu\text{mol/L}\cdot\text{h}^{-1}$  by CLA/ $\alpha$ -CD  
      microencapsules at 1:1, 1:2, 1:4, and 1:6 mol ratios, resp., indicating  
      that CLA oxidation was completely protected by a 1:4 mol ratio of  
      CLA/ $\alpha$ -CD. Such a protective effect by CLA/ $\beta$ -CD or  
      CLA/ $\gamma$ -CD microencapsules was achieved at a 1:6 mol ratio, but the  
      effect by CLA/ $\beta$ -CD was slightly greater than that by CLA/ $\gamma$ -CD.  
      The protective effect of  $\alpha$ -,  $\beta$ -, and  $\gamma$ -CDs for CLA oxidation  
      was confirmed by their POV-reducing abilities in CLA/CDs. These results  
      suggest that  $\alpha$ -CD was the most effective for the protection of CLA  
      oxidation by microencapsulation, followed by  $\beta$ -CD and  $\gamma$ -CD.  
AN    2000:554702    HCAPLUS <<LOGINID::20090302>>  
DN    133:265891  
TI    Improvement of Oxidative Stability of Conjugated Linoleic Acid  
      (CLA) by Microencapsulation in Cyclodextrins  
AU    Kim, Seck J.; Park, Gu B.; Kang, Chung B.; Park, Sang D.; Jung, Mun Y.;  
      Kim, Jeong O.; Ha, Yeong L.  
CS    Department of Agricultural Chemistry Animal Science and Veterinary  
      Medicine and Central Laboratory, Gyeongsang National University, Jinju,  
      660-701, S. Korea  
SO    Journal of Agricultural and Food Chemistry (2000), 48(9),  
      3922-3929  
      CODEN: JAFCAU; ISSN: 0021-8561  
PB    American Chemical Society  
DT    Journal  
LA    English

RE.CNT 45        THERE ARE 45 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8    ANSWER 7 OF 23    HCAPLUS    COPYRIGHT 2009 ACS on STN  
TI    Effects of amino acids, sugars, and ascorbic acid on the stability of  
      linoleic acid hydroperoxide in the water phase  
AB    Although lipid hydroperoxides are known to decrease food quality and  
      safety, the stability of hydroperoxides in foods has hardly been  
      investigated. Linoleic acid hydroperoxide (HPOD) decomposition by  
      kinetic means with or without various food components was examined. Most  
      amino acids, especially lysine, arginine and tryptophan, stabilized HPOD, while  
      cysteine and ascorbic acid accelerated its decomposition. Sugars had little  
      effect. According to activation energy calcns., it was found that the  
      HPOD decomposition mechanism in reaction systems with various food components  
      was similar to that in water.  
AN    1999:784767    HCAPLUS <<LOGINID::20090302>>  
DN    132:121634  
TI    Effects of amino acids, sugars, and ascorbic acid on the stability of  
      linoleic acid hydroperoxide in the water phase  
AU    Nishiike, Tamako; Ichikawa, Jun; Kikugawa, Noriko; Takamura, Hitoshi;  
      Matoba, Teruyoshi  
CS    Division of Human Life and Environmental Sciences, Graduate School of  
      Human Culture, Nara Women's University, Nara, 630-8506, Japan  
SO    Bioscience, Biotechnology, and Biochemistry (1999), 63(11),  
      1997-2000

CODEN: BBBIEJ; ISSN: 0916-8451

PB Japan Society for Bioscience, Biotechnology, and Agrochemistry  
DT Journal  
LA English

RE.CNT 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 8 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Oxidative stability and nuclear magnetic resonance analyses of  
linoleic acid encapsulated in cyclodextrins

AB The effects of  $\alpha$ - and  $\beta$ -cyclodextrin (CD) on the oxidative  
stability of linoleic acid (LA) at 35°C were studied by  
measuring headspace oxygen depletion in airtight 35-mL serum bottles. LA  
was encapsulated in  $\alpha$ -CD or  $\beta$ -CD in an aqueous solution during  
homogenization at 8000 rpm for 1 min and then dried under vacuum for 60 h  
at room temperature. Headspace oxygen was measured by thermal conductivity gas  
chromatog. The rate of oxygen depletion for the control, which contained  
LA only, was 93.8  $\mu$ mole/L·h. The rates of oxygen depletion for  
LA, encapsulated at a 1:1 mol ratio (mole CD/mol LA) in  $\alpha$ -CD and  
 $\beta$ -CD, were 13.8 and 111  $\mu$ moles/L·h, resp. When LA was  
encapsulated in  $\alpha$ -CD and  $\beta$ -CD at a 2:1 mol ratio (moles CD/mol  
LA), the rates of oxygen depletion were 0.573 and 53.9  
 $\mu$ moles/L·h, resp. Although  $\alpha$ -CD protected LA from  
reaction with oxygen at both ratios, the rate of oxygen depletion by LA  
encapsulated in  $\beta$ -CD at a 1:1 mol ratio was not statistically  
different from the control.  $\beta$ -CD protected LA from reaction with  
oxygen at a 2:1 mol ratio. 1H NMR spectra of the complexes formed from  
1:1 mol ratios of LA and CD indicated that LA was encapsulated in  
 $\alpha$ -CD or  $\beta$ -CD.

AN 1997:681639 HCAPLUS <<LOGINID::20090302>>

DN 127:358219

OREF 127:70123a,70126a

TI Oxidative stability and nuclear magnetic resonance analyses of  
linoleic acid encapsulated in cyclodextrins

AU Reichenbach, Wendy A.; Min, David B.

CS Department of Food Science, The Ohio State University, Columbus, OH,  
43210, USA

SO Journal of the American Oil Chemists' Society (1997), 74(10),  
1329-1333

CODEN: JAOCA7; ISSN: 0003-021X

PB AOCS Press

DT Journal

LA English

RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 9 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Mucosal preparation containing physiologically active peptide

AB This invention related to a mucosal preparation obtained by blending a physiol.  
active peptide at least with a sorbefacient and a vasodilatory compound.  
Owing to the combined use of the sorbefacient with the vasodilatory  
compound, the absorption of any desired physiol. active peptide can be  
enhanced and thus it can be self-administered to a patient without giving  
any pain caused by parenteral injection. Therefore, it is highly useful  
as a preparation of a physiol. active peptide for prolonged administration. As  
the physiol. active peptide, use can be made of insulin, calcitonin, human  
PTH, somatostatin, glucagon, etc. As the sorbefacient, use can be made of  
bile acid salts, cyclodextrin, phospholipids, nonionic surfactants, higher  
fatty acids, etc. As the vasodilatory compds., use can be made of calcium  
channel inhibitors, prostaglandin E1, isosorbide nitrate, nitroglycerin,  
etc.

AN 1997:259764 HCAPLUS <<LOGINID::20090302>>  
 DN 126:242891  
 OREF 126:46901a,46904a  
 TI Mucosal preparation containing physiologically active peptide  
 IN Yamamoto, Nakayuki; Ito, Teruomi  
 PA Asahi Kasei Kogyo Kabushiki Kaisha, Japan; Hisamitsu Seiyaku Kabushiki  
 Kaisha; Yamamoto, Nakayuki; Ito, Teruomi  
 SO PCT Int. Appl., 48 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA Japanese  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9706813	A1	19970227	WO 1996-JP2277	19960812 <--
	W: CA, CN, JP, KR, US				
	RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	JP 11292787	A	19991026	JP 1995-208010	19950815 <--
	CN 1179723	A	19980422	CN 1996-192821	19960812 <--
	EP 845265	A1	19980603	EP 1996-926626	19960812 <--
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
	JP 3824023	B2	20060920	JP 1997-509140	19960812 <--
PRAI	JP 1995-208010	A	19950815	<--	
	WO 1996-JP2277	W	19960812	<--	
OS	MARPAT 126:242891				

L8 ANSWER 10 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN  
 TI A method of producing a taxane-type diterpene  
 AB A simple method of producing a taxane-type diterpene by plant tissue culture is disclosed. Productivity can be improved by carrying out the culture in the presence of coronatines, a bacterium that produced the coronatines, a culture solution or a culture extract of such bacteria, cyclic polysaccharides, fatty acids, or an amino or imino derivative of jasmonic acids.

AN 1996:572123 HCAPLUS <<LOGINID::20090302>>  
 DN 125:219760  
 OREF 125:41103a,41106a  
 TI A method of producing a taxane-type diterpene  
 IN Yukimune, Yukihiro; Hara, Yasuhiro; Tan, Hiroaki; Tomino, Ikuro  
 PA Mitsui Petrochemical Industries, Ltd., Japan  
 SO Eur. Pat. Appl., 32 pp.  
 CODEN: EPXXDW

DT Patent  
 LA English

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 727492	A2	19960821	EP 1995-308498	19951127 <--
	EP 727492	A3	19961016		
	EP 727492	B1	20010131		
	R: DE, FR, GB, IT, NL				
	JP 08140690	A	19960604	JP 1994-291783	19941125 <--
	JP 3549594	B2	20040804		
	JP 08163991	A	19960625	JP 1994-312258	19941215 <--
	JP 09065889	A	19970311	JP 1995-218874	19950828 <--
	JP 3625908	B2	20050302		
	JP 08205882	A	19960813	JP 1995-301654	19951120 <--
	JP 3746550	B2	20060215		
PRAI	JP 1994-291783	A	19941125	<--	
	JP 1994-301179	A	19941205	<--	

JP 1994-312258 A 19941215 <--  
JP 1995-218874 A 19950828 <--  
OS MARPAT 125:219760

L8 ANSWER 11 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Aggregation of polyunsaturated fatty acids in the presence of cyclodextrins

AB The aggregation behavior of the polyunsatd. fatty acids (PUFA) linoleic acid and arachidonic acid was studied in the presence of cyclodextrins (CDs). The influence of CD concentration on CMC of PUFA suggests that two CD mols. bind sequentially to one mol. of PUFA. Two equilibrium consts., K1 representing the interaction of the first CD mol., and K2, the interaction of the second, were determined by non-linear regression of the PUFA CMC vs. CD concentration data to an expression deduced from the reaction scheme in the equilibrium. The effect of pH and the structure of the CD on the equilibrium

consts. was studied. It is postulated that the first CD mol. interacts with the carboxyl group of PUFA through hydrogen bonding when the fatty acid is protonated, while the second CD mol. binds to the hydrocarbon chain of the PUFA through hydrophobic interaction. The formation of hydrogen bonds was principally affected by the inner diameter of the CD, while the hydrophobic interactions were very strongly affected by the polarity of the CD group coating the inner channel. The relevance of the results for the development of enzyme assays involving fatty acids is discussed.

AN 1995:628687 HCAPLUS <<LOGINID::20090302>>

DN 123:50376

OREF 123:8923a,8926a

TI Aggregation of polyunsaturated fatty acids in the presence of cyclodextrins

AU Bru, Roque; Lopez-Nicolas, Jose M.; Garcia-Carmona, Francisco

CS Dep. Bioquim. Biol. Mol. "A", Univ. Murcia, Murcia, E-30001, Spain

SO Colloids and Surfaces, A: Physicochemical and Engineering Aspects ( 1995), 97(3), 263-9

CODEN: CPEAEH; ISSN: 0927-7757

PB Elsevier

DT Journal

LA English

L8 ANSWER 12 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Entrapment of liquid lipids into powdery matrixes of saccharides and proteins

AB The emulsifying activity, the high stabilizing activity of the emulsion and the formation of a fine dense skin layer during drying were the properties of agents that effectively entrapped liquid lipids. Gum arabic and gelatin were effective. Addition of an agent having a property to a base agent lacking the property improved the entrapment. Oxidation of entrapped liquid lipid was retarded. However, the extent of retardation depended on the kind of lipids and the kind of entrapping agents. Oxidation processes of some combinations of lipids and entrapping agents were expressed by a kinetic model including oxygen diffusion through dehydrated entrapping agents. Et eicosapentaenoate was also stabilized by the entrapment.

AN 1995:485889 HCAPLUS <<LOGINID::20090302>>

DN 122:263834

OREF 122:48177a,48180a

TI Entrapment of liquid lipids into powdery matrixes of saccharides and proteins

AU Matsuno, Ryuichi; Imagi, Jun; Adachi, Shuji

CS Fac. Agric., Kyoto Univ., Kyoto, 606-01, Japan

SO Dev. Food Eng., Proc. Int. Congr. Eng. Food, 6th (1994), Meeting

Date 1993, Volume Pt. 2, 1065-7. Editor(s): Yano, Toshimasa; Matsuno,

Ruuichi; Nakamura, Kozo. Publisher: Blackie, Glasgow, UK.  
CODEN: 61FFAL

DT Conference  
LA English

L8 ANSWER 13 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Utilization of cyclodextrin as fat soluble compound carrier to serum-free culture of rat astrocytes

AB  $\alpha$ -Cyclodextrin complexes with fat-soluble vitamins and unsatd. fatty acids were prepared and examined as replacements for bovine serum albumin as fat-soluble compound carriers on cultured rat astrocytes. In serum-supplemented medium, it was difficult to evaluate the effects of fat-soluble compds. in serum on cell growth. Therefore, serum-free chemical defined medium supplemented with growth factors, hormones, and nutrients was developed for rat astrocytes to evaluate these effects. .  
 $\alpha$ -Cyclodextrin complexes with 3 vitamins (vitamin A acetate, E, and K1) and 3 fatty acids (linoleic, linolenic, and oleic acids) showed growth promoting activities for astrocytes in serum-free medium. Usually, supplementing fat-soluble compds. to a cell culture medium is very difficult, especially to a low or no protein medium, but  $\alpha$ -cyclodextrin can replace albumin as a fat-soluble compound carrier in serum-free cell cultures.

AN 1993:579303 HCAPLUS <<LOGINID::20090302>>

DN 119:179303

OREF 119:32055a,32058a

TI Utilization of cyclodextrin as fat soluble compound carrier to serum-free culture of rat astrocytes

AU Nakama, Akihiko

CS Osaka City Inst. Public Health Environ. Sci., Osaka, 543, Japan

SO Annual Report of Osaka City Institute of Public Health and Environmental Sciences (1992), Volume Date 1991, 54, 48-53  
CODEN: AOISDR; ISSN: 0285-5801

DT Journal  
LA Japanese

L8 ANSWER 14 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Retarded oxidation of liquid lipids entrapped in matrixes of saccharides or proteins

AB Me linoleate (ML), linoleic acid (LA), and Et eicosapentaenoate (EE) were entrapped in saccharide and protein matrixes, and then stored at 37° in a desiccator controlled at 75% relative humidity. ML entrapped with  $\alpha$ -cyclodextrin, maltodextrin, and pullulan was extremely resistant to autoxidn., but LA entrapped with maltodextrin and pullulan rapidly oxidized. LA entrapped with  $\alpha$ -cyclodextrin was the most stable against oxidation. ML entrapped with gelatin or gum arabic was less resistant to autoxidn. than that entrapped with pullulan; there was little difference in the susceptibility to oxidation between ML and LA entrapped with gelatin or gum arabic. Egg albumin protected ML more effectively against oxidation than LA, while sodium caseinate protected LA more than ML. EE entrapped with pullulan was highly resistant to oxidation, 90% of the total lipid remaining after 35 days. The effect on the oxidation of diffusion of oxygen through the matrix was estimated. Retardation of oxidation of the entrapped lipid can

not

be explained only by the effect of diffusion.

AN 1992:590442 HCAPLUS <<LOGINID::20090302>>

DN 117:190442

OREF 117:32869a,32872a

TI Retarded oxidation of liquid lipids entrapped in matrixes of saccharides or proteins

AU Imagi, Jun; Muraya, Koji; Yamashita, Daisuke; Adachi, Shuji; Matsuno,

Ryuichi  
 CS Fac. Agric., Kyoto Univ., Kyoto, 606-01, Japan  
 SO Bioscience, Biotechnology, and Biochemistry (1992), 56(8),  
 1236-40  
 CODEN: BBBIEJ; ISSN: 0916-8451  
 DT Journal  
 LA English

L8 ANSWER 15 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN  
 TI Powderization of liquid-state lipids  
 AB Liquid-state lipids (linoleic acid, Me linoleate, or Me oleate)  
 were powderized by adsorption on gum arabic, starch, maltodextrin, .  
 alpha.-cyclodextrin, maltose, glucose, or CM-cellulose.  
 Lipids adsorbed on  $\alpha$ -cyclodextrin, gum arabic,  
 or CM-cellulose had high stability. The emulsifying activity of the  
 lipid-adsorbent complex is described.  
 AN 1991:654556 HCAPLUS <<LOGINID::20090302>>  
 DN 115:254556  
 OREF 115:43273a,43276a  
 TI Powderization of liquid-state lipids  
 AU Matsuno, Ryoichi; Imagi, Jun  
 CS Agric. Coll., Kyoto Univ., Kyoto, Japan  
 SO New Food Industry (1991), 33(5), 57-64  
 CODEN: NYFIAM; ISSN: 0547-0277  
 DT Journal  
 LA Japanese

L8 ANSWER 16 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN  
 TI Specific adsorbents in isolation and purification of cyclodextrins  
 AB A number of synthesized affinity sorbents were tested to find methods for the  
 separation of  $\alpha$ -,  $\beta$ -, and  $\gamma$ -cyclodextrins (CDs) from one  
 another and from acyclic dextrans. None of the gels retarded acyclic  
 dextrans, whereas  $\alpha$ -CD was specifically adsorbed onto supports  
 derivatized with alkyl functions,  $\beta$ -CD was specifically adsorbed onto  
 supports derivatized with phenyl or substituted Ph, and  $\gamma$ -CD was  
 specifically adsorbed onto a gel derivatized with a naphthyl compound. It  
 was evident that for achievement of binding capacities high enough for  
 practical preparation of the CDs, various parameters such as the support  
 material, its porosity, ligand, ligand concentration, temperature, and the  
 composition of  
 the mobile phase must be optimized.  
 AN 1989:453519 HCAPLUS <<LOGINID::20090302>>  
 DN 111:53519  
 OREF 111:9029a,9032a  
 TI Specific adsorbents in isolation and purification of cyclodextrins  
 AU Makela, Mauri; Mattsson, Pekka; Korpela, Timo  
 CS Dep. Biochem., Univ. Turku, Turku, SF-20500, Finland  
 SO Biotechnology and Applied Biochemistry (1989), 11(2), 193-200  
 CODEN: BABIEC; ISSN: 0885-4513  
 DT Journal  
 LA English

L8 ANSWER 17 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN  
 TI Effects of arachidonic acid and other long-chain fatty acids on the  
 membrane currents in the squid giant axon  
 AB The effects of arachidonic acid (I) and some other long-chain fatty acids  
 on the ionic currents of the voltage-clamped squid giant axon were  
 investigated by using intracellular application of the test substances.  
 The effects of these acids, which are usually insol. in solution, were examined  
 by using  $\alpha$ -cyclodextrin as a solvent. .  
 alpha.-Cyclodextrin itself had no effect on the

excitable membrane. I mainly suppressed the Na<sup>+</sup> current, but had little effect on the K<sup>+</sup> current. These effects were completely reversed after washing with control solution. The concentration required to suppress the peak inward current by 50% (ED<sub>50</sub>) was 0.18 mM, which was 10-fold larger than that of medium-chain fatty acids, like 2-decenoic acid. The Hill number was 1.5 for I, which is almost the same value as that for medium-chain fatty acids. This means that the mechanisms of the inhibition are similar in both long- and medium-chain fatty acids. When the long-chain fatty acids were compared, the efficacy of suppression of the Na<sup>+</sup> current was about the same value for I, docosatetraenoic acid, and docosahexaenoic acid. The suppression effects of linoleic acid and linolenic acid on Na<sup>+</sup> currents were 1/3 of that of I. Oleic acid had a small suppression effect and stearic acid had almost no effect on the Na<sup>+</sup> current. The currents were fitted to equations similar to those proposed by Hodgkin and Huxley (Hodgkin, A. L.; and Huxley, A. F., 1952) and the change in the parameters of these equations in the presence of fatty acids were calculated. The curve of the steady-state activation parameter for the Na<sup>+</sup> current against membrane potential and the time constant of activation were shifted 10 mV in a depolarizing direction by the application of fatty acids. The time constant for inactivation was almost unaffected by application of these fatty acids.

AN 1989:132753 HCAPLUS <<LOGINID::20090302>>

DN 110:132753

OREF 110:21875a,21878a

TI Effects of arachidonic acid and other long-chain fatty acids on the membrane currents in the squid giant axon

AU Takenaka, Toshifumi; Horie, Hidenori; Hori, Hideaki; Kawakami, Tadashi

CS Sch. Med., Yokohama City Univ., Yokohama, 236, Japan

SO Journal of Membrane Biology (1988), 106(2), 141-7

CODEN: JMBBBO; ISSN: 0022-2631

DT Journal

LA English

L8 ANSWER 18 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Growth of an established line of mouse mammary tumor cells under serum-free conditions

AB An established line of mouse mammary tumor cells (MTD cells) were cultured in a serum-free medium consisting of a 1:1 mixture of Dulbecco's modified Eagle's medium and Ham's F-12 medium supplemented with bovine serum albumin (BSA), insulin, and transferrin. To promote cell attachment and spreading, culture dishes were precoated with plasma fibronectin isolated from fibrinogen. Under these serum-free conditions, MTD cells grew at a rate close to that attained by the serum-supplemented medium. Among the additives in the serum-free medium, BSA was replaced with oleic acid or a complex of oleic acid and  $\alpha$ -cyclodextrin.

Transferrin was replaced with Fe<sup>2+</sup> or Fe<sup>3+</sup>. Addition of polyvinylpyrrolidone further improved the growth. Thus, MTD cells can be grown on a fibronectin-coated surface in a chemical defined medium with insulin as the only protein supplement. MTD cells grown under the serum-free conditions still retained the differentiated properties of the original MTD cells; i.e., the production of mouse mammary tumor virus in response to dexamethasone.

AN 1986:164689 HCAPLUS <<LOGINID::20090302>>

DN 104:164689

OREF 104:25993a,25996a

TI Growth of an established line of mouse mammary tumor cells under serum-free conditions

AU Kawamura, Kazuo; Enami, Jumpei; Kohmoto, Kaoru; Koga, Mutuyosi

CS Sch. Med., Dokkyo Univ., Mibu, 321-02, Japan

SO Dokkyo Journal of Medical Sciences (1985), 12(2), 167-80

CODEN: DJMSDB; ISSN: 0385-5023

DT Journal  
LA English

L8 ANSWER 19 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Medium for animal tissue culture

AB A medium containing cyclodextrin, to save serum, is prepared for animal tissue culture. Thus, 10 mg of linoleic acid, oleic acid, or vitamin E in 7 mL EtOH was added to an  $\alpha$ -cyclodextrin [10016-20-3] solution (1 g in 7 mL H<sub>2</sub>O) and heated to 70° under N<sub>2</sub>. When the solution turned transparent, it was rapidly cooled and kept cold (4°) for 20 h. The resulting precipitate was washed with 10 mL EtOH, dried under vacuum, washed with petroleum ether, and dried under vacuum. The RITC 56-2 medium was mixed with the 300 mg/L reaction product and 1 g/L  $\alpha$ -cyclodextrin and the ultrafiltered. Human lymphogemmul-like cells, UMCL-3, were cultured in the medium to yield .apprx.4.5 + 103 units of interferon/mL.

AN 1983:124208 HCAPLUS <<LOGINID::20090302>>

DN 98:124208

OREF 98:18913a,18916a

TI Medium for animal tissue culture

PA Ajinomoto Co., Inc., Japan; Yamane, Isao

SO Jpn. Kokai Tokkyo Koho, 11 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	---	-----	-----	-----
PI	JP 57194787	A	19821130	JP 1981-81600	19810528 <--
	JP 63018465	B	19880419		
PRAI	JP 1981-81600		19810528	<--	
OS	MARPAT 98:124208				

L8 ANSWER 20 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI  $\alpha$ -Cyclodextrin: a partial substitute for bovine serum albumin in serum-free culture of mammalian cells

AB The use was investigated of oleic acid- or linoleic acid- $\alpha$ -cyclodextrin inclusion complexes as albumin substitutes for mammalian cells.  $\alpha$ -Cyclodextrin did not show any cytotoxic effects at 2g/L medium. Growth curves are shown for 2 types of cells. UMCL cells grew well enough in the cyclodextrin-complex-containing, serum-free medium, whereas HEL cells required a small amount of albumin in addition to cyclodextrin for abundant growth.

AN 1982:612006 HCAPLUS <<LOGINID::20090302>>

DN 97:212006

OREF 97:35533a,35536a

TI  $\alpha$ -Cyclodextrin: a partial substitute for bovine serum albumin in serum-free culture of mammalian cells

AU Yamane, Isao; Kan, M.; Minamoto, Y.; Amatsuji, Y.

CS Inst. Tuberculosis Cancer, Tohoku Univ., Sendai, 980, Japan

SO Cold Spring Harbor Conferences on Cell Proliferation (1982), 9(Growth Cells Horm. Defined Media, Book A), 87-92

CODEN: CSHCAL; ISSN: 0097-5230

DT Journal

LA English

L8 ANSWER 21 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI  $\alpha$ -Cyclodextrin, a novel substitute for bovine albumin in serum-free culture of mammalian cells

AB The use of  $\alpha$ -,  $\beta$ -, and  $\gamma$ -cyclodextrin (CD) in combination with unsatd. fatty acids as a serum substitute in mammalian cell cultures



was examined by using a human lymphoblast cell line (UMCL) grown in RITC 56-1 medium supplemented with synthetic lecithin, cholesterol, galactose, and mannose and by using human diploid fibroblasts (HEL) grown in RITC 80-7 medium. On the basis of cytotoxic and cost considerations,  $\alpha$ -CD was used for the expts. Both  $\alpha$ -CD-oleic acid and  $\alpha$ -CD-linoleic acid had growth-enhancing effects on UMCL cells up to 100 mg/L medium but exhibited toxic effects at higher concns. However, when 100 mg  $\alpha$ -CD included with both fatty acids and 1000 mg free  $\alpha$ -CD were added to 1 L of medium, stable and reproducible growth-promoting effects were observed. With HEL cells, growth similar to that in bovine serum albumin-supplemented medium was observed by addition of a concentrated  $\alpha$ -CD complex to a final concentration of 10-20 mg/L.

AN 1982:100488 HCAPLUS <<LOGINID::20090302>>

DN 96:100488

OREF 96:16453a,16456a

TI  $\alpha$ -Cyclodextrin, a novel substitute for bovine albumin in serum-free culture of mammalian cells

AU Yamane, Isao; Kan, Mikio; Minamoto, Yoshiki; Amatsuji, Yasuo

CS Res. Inst. Tuberc. Cancer, Tohoku Univ., Sendai, 980, Japan

SO Proceedings of the Japan Academy, Series B: Physical and Biological Sciences (1981), 57(10), 385-9  
CODEN: PJABDW; ISSN: 0386-2208

DT Journal

LA English

L8 ANSWER 22 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Chromatographic investigation of the component glycerides of kusum [kusam] oil

AB Sep. the triglyceride components from a 5% solution of crude kusum or Macassar oil (from the seeds of *Schleichera trijuga*) in CHCl<sub>3</sub> by adsorption thin-layer chromatog. (t.l.c.) on Silica Gel G plates by developing with the 75:25:1 n-C<sub>6</sub>H<sub>14</sub>-Et<sub>2</sub>O-HOAc solvent of D. C. Malins and H. K. Mangold (1960). Locate the bands by exposure to I, remove adsorbed I, extract the adsorbent with warm CHCl<sub>3</sub>, and filter. Concentrate the filtrate to

a 1% solution by evaporation. Subfractionate the triglyceride components by t.l.c.

on Silica Gel G plates impregnated with AgNO<sub>3</sub>, by developing with the 100:0.5 CHCl<sub>3</sub>-HOAc solvent of C. B. Barrett, et al. (1963). Locate the bands by spraying with 0.2% 2',7'-dichlorofluorescein in 95% EtOH and exposing to uv light. Extract the bands with anhydrous Et<sub>2</sub>O and free from the dye by the procedure of H. P. Kaufmann and H. Wessels (1966). Evaporate the solvent from each fraction, then saponify with N KOH in EtOH, remove EtOH, acidify to liberate the fatty acids, extract with Et<sub>2</sub>O, and wash free of mineral acids with H<sub>2</sub>O. Remove the Et<sub>2</sub>O and dissolve each residual fraction in 0.5 ml. CHCl<sub>3</sub>. Fractionate the fatty acids of each fraction by reversed-phase t.l.c. on a plaster of Paris plate coated with 5% liquid paraffin in petroleum ether, as described by H. P. Kaufmann, et al. (1961). Develop the plates with 90% HOAc or 70% HOAc for the higher- and lower-mol.-weight fatty acids, resp. Detect the spots by spraying with 1%  $\alpha$ -cyclodextrin in 30% EtOH and exposing to iodine. The fatty acid compns. of the 10 triglyceride subfractions obtained by AgNO<sub>3</sub> t.l.c. are presented.

AN 1969:431621 HCAPLUS <<LOGINID::20090302>>

DN 71:31621

OREF 71:5853a,5856a

TI Chromatographic investigation of the component glycerides of kusum [kusam] oil

AU Kundu, M. K.

CS Calcutta Univ., Calcutta, India

SO Journal of Chromatography (1969), 41(2), 276-8

CODEN: JOCRAM; ISSN: 0021-9673

DT Journal  
LA English

L8 ANSWER 23 OF 23 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Acylated cyclodextrins as polar stationary phases for gas-liquid chromatography

AB The use of  $\beta$ -cyclodextrin acetate (I) (mol. weight 2018, m. 199-201°), and of mixts. of I,  $\beta$ -cyclodextrin propionate (mol. weight 2312, m. 169°), and  $\alpha$ -cyclodextrin acetate (mol. weight 1730, m. 243-5°) as stationary phases for gas-liquid chromatography is recommended. Very little bleeding or degradation was evident for 10 g. I applied on 40 g. 30-60 mesh Chromasorb R, when used at 236° with a 57 ml./min. He flow through a 10 ft. + 1/4-in. column. Resolution of fatty acids obtained with the above column resembled that obtained with a butanediol-succinic acid polyester stationary phase under similar conditions.

AN 1962:10665 HCAPLUS <<LOGINID::20090302>>

DN 56:10665

OREF 56:1983a-c

TI Acylated cyclodextrins as polar stationary phases for gas-liquid chromatography

AU Sand, Donald M.; Schlenk, Hermann

CS Univ. of Minnesota, Austin

SO Anal. Chem. (1961), 33, 1624-5

CODEN: ANCHAM; ISSN: 0003-2700

DT Journal

LA Unavailable